Declaration of Conformity

CERTIFICATE & DECLARATION OF CONFORMITY FOR CE MARKING

The above medical devices are classified according to the regulation (EU) MDR 2017/745 for medical devices.

Manufacturer's name:	Vigor Dongguan Factory
Attn.	Mr. Forrest Feng
Business address:	The 5th Floor, Building 2, Baili Industrial Park, No.382 Liaodong Road, Xixi Village, Liaobu Town, Dongguan City, Guangdong Province, China.
SRN number:	CN-MF-000021497
Basic UDI-ID:	697368741
EU Representative:	Nygård Jensen APS (Vigor Europe), Havnevej 1, 4000 Roskilde, Denmark.
SRN number:	DK-AR-000019275
Basic UDI-ID:	697368741
Medical device(s):	**Medical Slings (Patient lifting Slings)**
Classification:	Class I
Scope of application:	OEM, ODM manufactured Slings.

The above medical devices are classified according to the regulation (EU) MDR 2017/745 for medical devices.

The products comply with REACH regulations and do not contain PVC/Cadmium.

The conformity assessment and verification procedure are supported by the Quality System approval issued by NQA UKAS.

It is certified that all slings are checked and tested according to ISO 10535:2021.

ISO-13485:2016 Certificate number: 131095

Authorised signatory:

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Signature

Forrest Feng, General Manager

Date: 7-11-2024

CE

This declaration of conformity is issued by Vigor Dongguan Factory, and we hereby declare that the medical device(s) specified above meet the provision of the Regulation (EU) MDR 2017/745 for medical devices. All supporting documentation is retained at the premises of the manufacturer.